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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/518,297	03/03/2000	Moon Young Lim	4600-0130.30 5390	
22918 7.	590 03/31/2006		EXAMINER	
PERKINS COIE LLP			KAM, CHIH MIN	
P.O. BOX 2168 MENLO PARK, CA 94026			ART UNIT	PAPER NUMBER
•			1656	
			DATE MAILED: 03/31/2006	

Please find below and/or attached an Office communication concerning this application or proceeding.

<del></del>	Application No.	L A II A/ . \
	Application No.	Applicant(s)
Office Action Summer	09/518,297	LIM ET AL.
Office Action Summary	Examiner	Art Unit
	Chih-Min Kam	1656
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with the o	correspondence address
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D.  - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period.  - Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICATION  136(a). In no event, however, may a reply be tir  will apply and will expire SIX (6) MONTHS from  e, cause the application to become ABANDONE	N. nely filed the mailing date of this communication. ED (35 U.S.C. § 133).
Status		
1) ■ Responsive to communication(s) filed on <u>05 C</u> 2a) ■ This action is <b>FINAL</b> . 2b) ■ This      3) ■ Since this application is in condition for alloware closed in accordance with the practice under the practice.	s action is non-final. ince except for formal matters, pro	
Disposition of Claims		
4)  Claim(s) 52,53 and 60-63 is/are pending in the 4a) Of the above claim(s) is/are withdra 5)  Claim(s) 52 and 53 is/are allowed.  6)  Claim(s) 60-63 is/are rejected.  7)  Claim(s) is/are objected to.  8)  Claim(s) are subject to restriction and/o	wn from consideration.	
Application Papers		
9)☑ The specification is objected to by the Examine 10)☐ The drawing(s) filed on is/are: a)☐ acc Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11)☐ The oath or declaration is objected to by the Examine 11.	cepted or b) objected to by the drawing(s) be held in abeyance. Set tion is required if the drawing(s) is objected to by the drawing(s).	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).
Priority under 35 U.S.C. § 119		
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. Its have been received in Applicationity documents have been received u (PCT Rule 17.2(a)).	on No ed in this National Stage
Attachment(s)		
Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal P 6) Other:	(PTO-413) ate atent Application (PTO-152)

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#### **DETAILED ACTION**

1. The instant application has been withdrawn from issue (see the notice dated February 1, 2006), and an office action follows.

### Status of the Claims

2. Claims 52, 53 and 60-63 are pending.

Applicants' amendment to drawings filed October 5, 2005 is acknowledged. Claims 52, 53 and 60-63 are examined.

## Withdrawn Objection to Drawings

3. The previous objection of to Figs. 2A and 2B is withdrawn in view of applicants' submission of formal drawings in the amendment filed October 5, 2005.

# Objection to New Matter Added to Specification

4. The preliminary amendment filed May 22, 2000 and amendment filed May 12, 2005 are objected to under 35 U.S.C. 132 because they introduce new matter into the disclosure. 35 U.S.C. 132 states that no amendment shall introduce new matter into the disclosure of the invention. The added material which is not supported by the original disclosure is as follows: The original specification does not disclose

TAPITDVSLGDELRLDGEEVDMTPMMDDFDLEMLGDVESPSPGMTHDPVSYGMD VDDFEFEQMFTDALGIDDFG as SEQ ID NO:8 and

ADALDDFDLEMADALDDFDLEMADALDDFDLEM as SEQ ID NO:9 in the Sequence Listing Table (see amendment filed May 12, 2005); and the amino acid sequences of SEQ ID NO:63 and 64 (for herpes simplex virus type 2 V16 Genbank Accession number

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M57289) in the Sequence Listing Table (see amendment filed May 12, 2005), and Sequence Listing.

Applicant is required to cancel the new matter in the reply to this Office Action.

## Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

4. Claims 60-63 are rejected under 35 U.S.C. 102(b) as being anticipated by Bujard *et al.* (WO 94/29442).

Bujard *et al.* teach a system for regulating expression of eucaryotic genes using components of the Tet repressor/operator/inducer system of prokaryotes in a host cell, and transcription of a nucleotide sequence operably linked to at least one tet operator sequence is stimulated by a tetracycline (Tc)-controllable transcriptional activator fusion protein (tTA) which comprises two polypeptides, the first polypeptide is a Tet repressor (TetR), which binds to tet operator sequence in the absence of Tc, and the second polypeptide directly or indirectly activates transcription in eucaryotic cells, e.g., the second polypeptide can be a transcriptional activation domain from herpes simplex virus viron protein 16 (VP16) (page 2, lines 4-14), where in the absence of Tc, transcription of a gene operably linked to a tTA-responsive promoter (typically comprising at least one tet operator sequence and a minimal promoter) is stimulated by a tTA (page 2, lines 17-21). The reference also indicates a host cell can contain a polynucleotide moiety encoding a tTA and a gene of interest operably linked to a tTA-responsive transcriptional

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promoter, where the gene of interest operably linked to the tTA-responsive transcriptional promoter can be integrated into DNA of the host cell either randomly (e.g., by introduction of an exogenous gene) or at a predetermined location (e.g., by targeting an endogenous gene for homologous recombination, the integration of polynucleotide encoding tTA and a tTA-responsive promoter, page 3, lines 24-39; claims 60-63), and expression of a gene of interest operably linked to a tTA-responsive transcriptional promoter in a host cell can be inhibited by contacting the cell with Tc (page 4, line 1-31; page 12, lines 19-37). The tTA in the absence of Tc is a DNA binding compound and modulates the binding of the transcriptional activation domain such as VP16 to the DNA response element, which meets the criteria of the claims.

Perhaps claims 60-63 can be amended as follows to differentiate from the prior art:

- 60. A molecular switch, comprising:
- (a) a first nucleic acid construct, having
- (i) a DNA response element for a transcriptional regulatory protein, operably linked to a first promoter;
- (ii) a non-native compound binding sequence which is the same as, overlapping, or adjacent to said DNA response element, for binding to a DNA binding compound;
  - (iii) a transgene under the control of said first promoter; and
- (b) a DNA binding compound, wherein the DNA binding compound is separate and different from the transcriptional regulatory protein; and
- (c) a second nucleic acid construct, having the coding sequence for said transcriptional regulatory protein operably linked to a second promoter;

wherein said DNA binding compound, when bound to said binding sequence, is effective to modulate binding of said transcriptional regulatory protein to said DNA response element and wherein a first vector is including said first nucleic acid construct and a second vector is including said second nucleic acid construct.

- 61. A molecular switch, comprising:
- (a) a first nucleic acid construct, having
- (i) a DNA response element for a transcriptional regulatory protein, operably linked to a first promoter;

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(ii) a non-native compound binding sequence which is the same as, overlapping, or adjacent to said DNA response element, for binding to a DNA binding compound;

- (iii) a transgene under the control of said first promoter; and
- (b) a DNA binding compound, wherein the DNA binding compound is separate and different from the transcriptional regulatory protein;

wherein said DNA binding compound, when bound to said binding sequence, is effective to modulate binding of said transcriptional regulatory protein to said DNA response element and wherein said compound binding sequence has from about 8 to 20 nucleotides.

- 62. A molecular switch, comprising:
- (a) a first nucleic acid construct, having
- (i) a DNA response element for a transcriptional regulatory protein, operably linked to a first promoter;
- (ii) a non-native compound binding sequence which is the same as, overlapping, or adjacent to said DNA response element, for binding to a DNA binding compound;
  - (iii) a transgene under the control of said first promoter; and
- (b) a DNA binding compound, wherein the DNA binding compound is separate and different from the transcriptional regulatory protein;

wherein said DNA binding compound, when bound to said binding sequence, is effective to modulate binding of said transcriptional regulatory protein to said DNA response element and wherein said nucleic acid construct has from 1 to 12 compound binding sequences.

- 63. A molecular switch, comprising:
- (a) a first nucleic acid construct, having
- (i) a DNA response element for a transcriptional regulatory protein, operably linked to a first promoter;
- (ii) a non-native compound binding sequence which is the same as, overlapping, or adjacent to said DNA response element, for binding to a DNA binding compound;
  - (iii) a transgene under the control of said first promoter; and
- (b) a DNA binding compound, wherein the DNA binding compound is separate and different from the transcriptional regulatory protein;

wherein said DNA binding compound, when bound to said binding sequence, is effective to modulate binding of said transcriptional regulatory protein to said DNA response element and wherein said nucleic acid construct has from 1 to 12 tandom repeated transcriptional regulatory protein DNA response elements.

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#### Conclusion

5. Claims 60-63 are rejected. It appears claims 52 and 53 are free of prior art.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Chih-Min Kam whose telephone number is (571) 272-0948. The examiner can normally be reached on 8.00-4:30, Mon-Fri.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Kathleen Kerr can be reached at 571-272-0931. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Chif

Chih-Min Kam, Ph. D.

Patent Examiner

CHIH-MIN KAM
PATENT EXAMINER

**CMK** 

March 24, 2006